Application Serial No.: 10/680,673 Amendment dated: February 7, 2007 Response to Office Action dated August 7, 2007

## Amendments to the Claims:

This listing of the claims will replace all prior versions, and listings, of claims in the application:

## Listing of Claims:

- (Withdrawn) A composition for intraarticular delivery of chondrogenic polypeptides comprising a pharmaceutically acceptable admixture comprising FGF18 and hyaluronic acid.
- 2. (Withdrawn) The composition of claim 1 further comprising a negatively charged carrier.
- 3. (Withdrawn) The composition of claim 2 wherein said carrier is selected from the group consisting of low molecular weight hyaluronans, sulfated proteoglycans, B-cyclodextrin tetradecasulphate, hydroxyapatite, alginate microspheres, chitosans, and methylcellulose.
- 4. (Withdrawn) The composition of claim 1 wherein said composition is a time-release formulation.
- (Withdrawn) The composition of claim 4 wherein said time-release formulation comprises a matrix selected from the group consisting of a solution, a gel, a paste, or a putty.
- (Withdrawn) The composition of claim 4 wherein said time-release formulation comprises a reservoir system.
- (Withdrawn) The composition of claim 1 further comprising chondrocytes wherein said chondrocytes have been cultured in the presence of FGF18 prior to intraarticular administration.
- 8. (Withdrawn) The composition of claim 1 further comprising an anti-inflammatory drug.

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- 9. (Presently amended) A method for increasing chondrocyte proliferation in a joint of a mammal in need thereof comprising the step of administering into a synovial cavity a pharmaceutically acceptable admixture comprising consisting of FGF18 and hyaluronic acid.
- 10. (Original) The method of claim 9 wherein said administration comprises injection.
- 11. (Original) The method of claim 9 wherein said administration comprises surgical implantation.
- 12. (Presently amended) The method of claim 9 wherein said admixture administration further comprises a negatively-charged carrier selected from the group consisting of low molecular weight hyaluronans, high molecular weight hyaluronans, sulfated proteoglycans, B-cyclodextrin tetradecasulphate, hydroxyapatite, alginate microspheres, chitosans, and methylcellulose.
- 13. (Presently amended) The method of claim 9 where said admixture is a time-release formulation administration occurs over time.
- 14. (Presently amended) The method of claim 13 wherein said administration over time is accomplished using time-release formulation-comprises a matrix selected from the group consisting of a solution, a gel, a paste, or a putty.
- 15. (Presently amended) The method of claim 13 wherein said administration over time is accomplished using time-release-formulation-comprises a reservoir system.
- (Presently amended) The method of claim 9 wherein said admixture administration further comprises an anti-inflammatory drug.
- 17. (Original) The method of claim 9 further comprising the steps of allowing growth of new cartilage tissue and surgically contouring the new cartilage surface.

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- 18. (Presently amended) A method of treating osteoarthritis in a mammal comprising the steps of administering into a synovial cavity a pharmaceutically acceptable admixture eomprising consisting of FGF18 and hyaluronic acid.
- (Original) The method of claim 18 wherein said administration comprises injection.
- (Original) The method of claim 18 wherein said administration comprises surgical implantation.
- 21. (Presently amended) The method of claim 18 wherein said admixture administration further comprises a negatively-charged carrier selected from the group consisting of low molecular weight hyaluronans, high molecular weight hyaluronans, sulfated proteoglycans, B-cyclodextrin tetradecasulphate, hydroxyapatite, alginate microspheres, chitosans, and methylcellulose.
- 22. (Presently amended) The method of claim 18 where said admixture is a time release formulation administration occurs over time.
- 23. (Presently amended) The method of claim 22 wherein said administration over time is accomplished using time release fermulation comprises a matrix selected from the group consisting of a solution, a gel, a paste, or a putty.
- 24. (Presently amended) The method of claim 22 wherein said administration over time is accomplished using time release formulation-comprises a reservoir system.
- 25. (Presently amended) The method of claim 18 wherein said admixture administration further comprises an anti-inflammatory drug.
- 26. (Original) The method of claim 18 further comprising the steps of allowing growth of new cartilage tissue and surgically contouring the new cartilage surface.